

V. Balasubramanian

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```
=> file reg
COST IN U.S. DOLLARS          SINCE FILE      TOTAL
                                ENTRY        SESSION
FULL ESTIMATED COST          0.21           0.21
```

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=>Testing the current file.... screen

ENTER SCREEN EXPRESSION OR (END):end

=>
Uploading C:\STNEXP4\QUERIES\099951592.str

L1 STRUCTURE uploaded

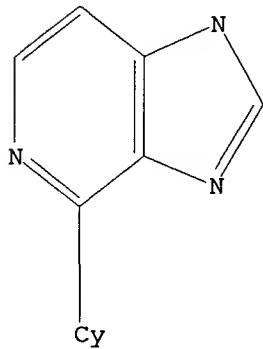
=> que L1

L2 QUE L1

=> d l1

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=>Testing the current file.... screen

ENTER SCREEN EXPRESSION OR (END):end

=>

Uploading C:\STNEXP4\QUERIES\099951591.str

L3 STRUCTURE UPLOADED

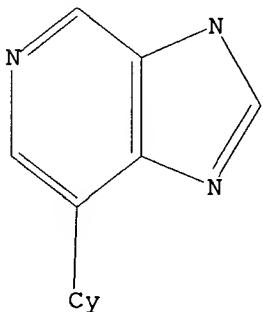
=> que L3

L4 QUE L3

=> d 13

L3 HAS NO ANSWERS

L3 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1 or l3 sss sam

SAMPLE SEARCH INITIATED 13:33:19 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 1457 TO ITERATE

68.6% PROCESSED 1000 ITERATIONS

0 ANSWERS

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INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.03

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 26851 TO 31429
PROJECTED ANSWERS: 0 TO 0

L5 0 SEA SSS SAM L1 OR L3

=> s l1 or l3 sss ful
FULL SEARCH INITIATED 13:33:38 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 29137 TO ITERATE

100.0% PROCESSED 29137 ITERATIONS 11 ANSWERS
SEARCH TIME: 00.00.05

L6 11 SEA SSS FUL L1 OR L3

=> file caplus
COST IN U.S. DOLLARS SINCE FILE TOTAL
FULL ESTIMATED COST ENTRY SESSION
214.54 214.75

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FILE COVERS 1907 - 12 Jul 2002 VOL 137 ISS 3
FILE LAST UPDATED: 11 Jul 2002 (20020711/ED)

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=> s 16
L7 5 L6

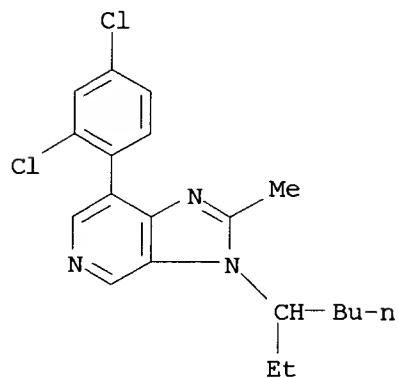
=> d 17 1-5 bib hitstr

L7 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS
AN 1998:568830 CAPLUS

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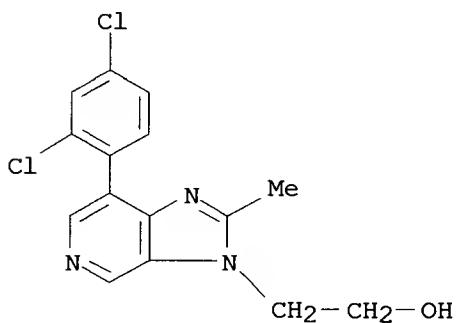
DN 129:202953
 TI Preparation of bicyclic nitrogen-containing heterocycles as CRF receptor antagonists and methods relating thereto
 IN McCarthy, James R.
 PA Neurocrine Biosciences, Inc., USA
 SO PCT Int. Appl., 62 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------|---|------|----------|-----------------|----------|
| PI | WO 9835967 | A2 | 19980820 | WO 1998-US2932 | 19980217 |
| | WO 9835967 | A3 | 19981210 | | |
| | W: AL, AM, AT, AU, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM | | | | |
| | RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG | | | | |
| | AU 9862795 | A1 | 19980908 | AU 1998-62795 | 19980217 |
| | EP 970082 | A2 | 20000112 | EP 1998-905094 | 19980217 |
| | R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI | | | | |
| | JP 2001511813 | T2 | 20010814 | JP 1998-535982 | 19980217 |
| | US 6352990 | B1 | 20020305 | US 1999-415503 | 19991008 |
| | US 2002049207 | A1 | 20020425 | US 2001-995159 | 20011127 |
| PRAI | US 1997-36414P | P | 19970218 | | |
| | US 1997-36415P | P | 19970218 | | |
| | US 1997-36416P | P | 19970218 | | |
| | US 1997-36421P | P | 19970218 | | |
| | US 1997-36422P | P | 19970218 | | |
| | US 1997-36423P | P | 19970218 | | |
| | WO 1998-US2932 | W | 19980217 | | |
| | US 1999-415503 | A3 | 19991008 | | |
| OS | MARPAT 129:202953 | | | | |
| IT | 212138-53-9P , 7-(2,4-Dichlorophenyl)-2-methyl-3-(hept-3-yl)-3H-imidazo[4,5-c]pyridine 212138-55-1P , 7-(2,4-Dichlorophenyl)-2-methyl-3-(2-hydroxyethyl)-3H-imidazo[4,5-c]pyridine | | | | |
| | RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) | | | | |
| | (prepn. of bicyclic nitrogen heterocycles as CRF receptor antagonists) | | | | |
| RN | 212138-53-9 CAPLUS | | | | |
| CN | 3H-Imidazo[4,5-c]pyridine, 7-(2,4-dichlorophenyl)-3-(1-ethylpentyl)-2-methyl- (9CI) (CA INDEX NAME) | | | | |



RN 212138-55-1 CAPLUS

CN 3H-Imidazo[4,5-c]pyridine-3-ethanol, 7-(2,4-dichlorophenyl)-2-methyl- (9CI) (CA INDEX NAME)



L7 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2002 ACS

AN 1997:706254 CAPLUS

DN 128:3662

TI Dienophilicity of imidazole in inverse electron demand Diels-Alder reactions; intermolecular reactions with 1,2,4-triazines

AU Wan, Zhaokui; Snyder, John K.

CS Department of Chemistry, Boston University, Boston, MA, 02215, USA

SO Tetrahedron Letters (1997), 38(43), 7495-7498

CODEN: TELEAY; ISSN: 0040-4039

PB Elsevier

DT Journal

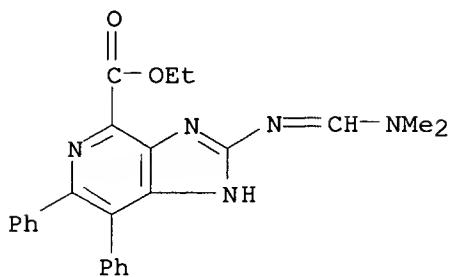
LA English

IT 198707-68-5P

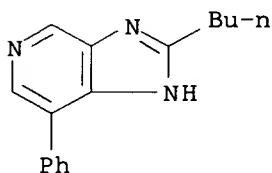
RL: SPN (Synthetic preparation); PREP (Preparation)
(Diels-Alder reactions of imidazoles with triazines)

RN 198707-68-5 CAPLUS

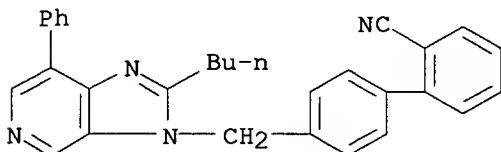
CN 1H-Imidazo[4,5-c]pyridine-4-carboxylic acid, 2-
[[(dimethylamino)methylene]amino]-6,7-diphenyl-, ethyl ester (9CI) (CA
INDEX NAME)



L7 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2002 ACS
 AN 1995:741580 CAPLUS
 DN 123:339884
 TI Synthesis and evaluation of novel nonpeptide angiotensin II receptor antagonists: imidazo[4,5-c]pyridine derivatives with an aromatic substituent
 AU Kiyama, Ryuichi; Fuji, Masahiro; Hara, Mariko; Fujimoto, Masafumi; Kawabata, Tomoji; Nakamura, Matsuhsia; Fujishita, Toshio
 CS Shionogi Res. Lab., Shionogi & Co., Ltd., Osaka, 553, Japan
 SO Chemical & Pharmaceutical Bulletin (1995), 43(3), 450-60
 CODEN: CPBTAL; ISSN: 0009-2363
 PB Pharmaceutical Society of Japan
 DT Journal
 LA English
 IT 170505-66-5P 170505-70-1P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (for prepn. of imidazo[4,5-c]pyridine derivs. with arom. substituent as antagonists)
 RN 170505-66-5 CAPLUS
 CN 1H-Imidazo[4,5-c]pyridine, 2-butyl-7-phenyl- (9CI) (CA INDEX NAME)



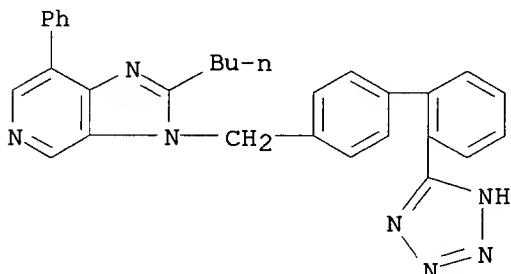
RN 170505-70-1 CAPLUS
 CN [1,1'-Biphenyl]-2-carbonitrile, 4'-[(2-butyl-7-phenyl-3H-imidazo[4,5-c]pyridin-3-yl)methyl]- (9CI) (CA INDEX NAME)



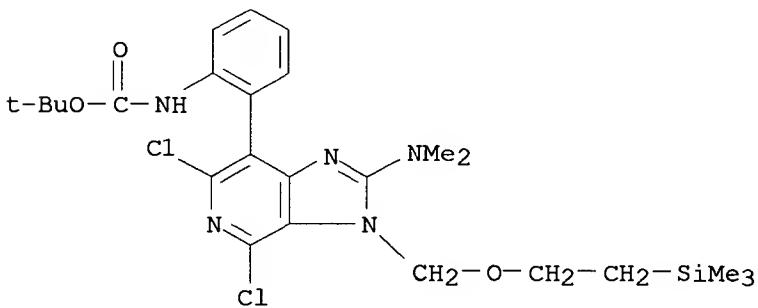
IT 170506-01-1P

V. Balasubramanian

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. as nonpeptide angiotensin II receptor antagonists)
 RN 170506-01-1 CAPLUS
 CN 3H-Imidazo[4,5-c]pyridine, 2-butyl-7-phenyl-3-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl)methyl]- (9CI) (CA INDEX NAME)



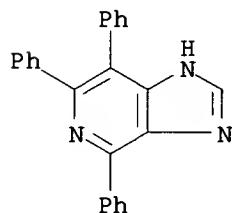
L7 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2002 ACS
 AN 1993:580583 CAPLUS
 DN 119:180583
 TI A short entry into the pyrido[2,3-b]indole ring system. Synthesis of the tetracyclic segment of the marine antitumor agents: grossularines-1 and -2
 AU Achab, Said; Guyot, Michele; Potier, Pierre
 CS Lab. Chim., Mus. Natl. Histor. Nat., Paris, F 75231, Fr.
 SO Tetrahedron Lett. (1993), 34(13), 2127-30
 CODEN: TELEAY; ISSN: 0040-4039
 DT Journal
 LA English
 OS CASREACT 119:180583
 IT **148493-57-6P**
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (prepn. and cyclization of)
 RN 148493-57-6 CAPLUS
 CN Carbamic acid, [2-[4,6-dichloro-2-(dimethylamino)-3-[[2-(trimethylsilyl)ethoxy]methyl]-3H-imidazo[4,5-c]pyridin-7-yl]phenyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



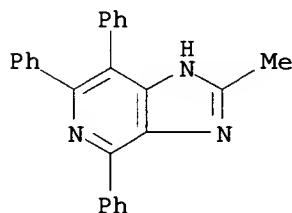
L7 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2002 ACS
 AN 1983:198113 CAPLUS

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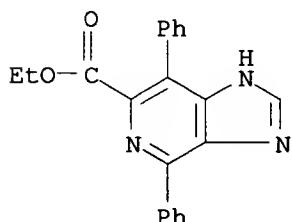
DN 98:198113
 TI Reduction of 4,7-diphenyl-1,2,5-thia(oxa)diazolo[3,4-c]pyridines affording 2,5-diphenyl-3,4-diaminopyridines and ring closure of the diamines to fluorescent azaheterocycles
 AU Mataka, Shuntaro; Takahashi, Kazufumi; Imura, Tetsuro; Tashiro, Masashi
 CS Res. Inst. Ind. Sci., Kyushu Univ. 86, Kasuga, 816, Japan
 SO J. Heterocycl. Chem. (1982), 19(6), 1481-8
 CODEN: JHTCAD; ISSN: 0022-152X
 DT Journal
 LA English
 OS CASREACT 98:198113
 IT **85731-48-2P 85731-49-3P 85731-50-6P**
85731-52-8P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (prepn. of)
 RN 85731-48-2 CAPLUS
 CN 1H-Imidazo[4,5-c]pyridine, 4,6,7-triphenyl- (9CI) (CA INDEX NAME)



RN 85731-49-3 CAPLUS
 CN 1H-Imidazo[4,5-c]pyridine, 2-methyl-4,6,7-triphenyl- (9CI) (CA INDEX NAME)

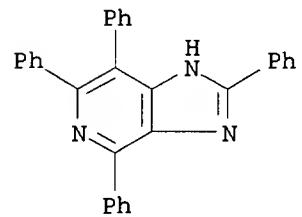


RN 85731-50-6 CAPLUS
 CN 1H-Imidazo[4,5-c]pyridine-6-carboxylic acid, 4,7-diphenyl-, ethyl ester (9CI) (CA INDEX NAME)



V. Balasubramanian

RN 85731-52-8 CAPLUS
CN 1H-Imidazo[4,5-c]pyridine, 2,4,6,7-tetraphenyl- (9CI) (CA INDEX NAME)



| => log y | | | |
|----------------------|------------|---------|--|
| COST IN U.S. DOLLARS | SINCE FILE | TOTAL | |
| FULL ESTIMATED COST | ENTRY | SESSION | |
| | 15.67 | 230.42 | |

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| NEWS 22 Jul 02 | FOREGE no longer contains STANDARDS file segment |
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| | | | |
|----------------------|--|------------|---------|
| => file reg | | SINCE FILE | TOTAL |
| COST IN U.S. DOLLARS | | ENTRY | SESSION |
| FULL ESTIMATED COST | | 0.21 | 0.21 |

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ENTER SCREEN EXPRESSION OR (END):end

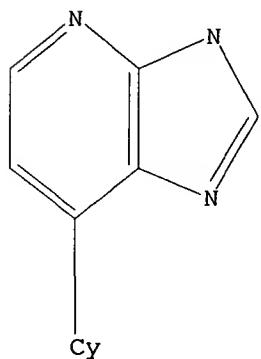
=>
Uploading C:\STNEXP4\QUERIES\099951593.str

L1 STRUCTURE uploaded

=> que L1

L2 QUE L1

=> d l1
L1 HAS NO ANSWERS
L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s ll sss sam

SAMPLE SEARCH INITIATED 13:27:07 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 1311 TO ITERATE

76.3% PROCESSED 1000 ITERATIONS 0 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 24049 TO 28391
PROJECTED ANSWERS: 0 TO 0

L3 0 SEA SSS SAM L1

=> s ll sss ful

FULL SEARCH INITIATED 13:27:26 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 25799 TO ITERATE

100.0% PROCESSED 25799 ITERATIONS 3 ANSWERS
SEARCH TIME: 00.00.04

L4 3 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS

| | SINCE FILE ENTRY | TOTAL SESSION |
|---------------------|------------------|---------------|
| FULL ESTIMATED COST | 140.66 | 140.87 |

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=> s 14
L5 3 L4

=> d 15 1-3 bib hitstr

L5 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2002 ACS
AN 2002:31438 CAPLUS
DN 136:102370
TI Preparation of tetrahydropyridine or piperidine heterocyclic derivatives and their affinity for CRF receptors
IN Nakazato, Atsuro; Kumagai, Toshihito; Okubo, Taketoshi; Kameo, Kazuya
PA Taisho Pharmaceutical Co., Ltd., Japan
SO PCT Int. Appl., 91 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

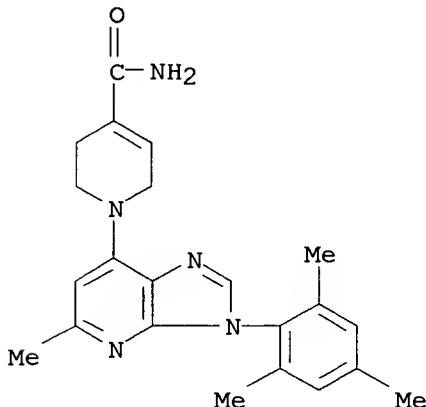
| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------|--|--|----------|--|--|
| PI | WO 2002002549 | A1 | 20020110 | WO 2001-JP5806 | 20010704 |
| | W: | AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM | RW: | GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG | AU 2001069437 A5 20020114 AU 2001-69437 20010704 |
| PRAI | JP 2000-204021 | A | 20000705 | | |
| | JP 2000-270535 | A | 20000906 | | |
| | JP 2000-2000204021A | | 20000705 | | |
| | JP 2000-2000270535A | | 20000906 | | |
| | WO 2001-JP5806 | W | 20010704 | | |
| OS | MARPAT | 136:102370 | | | |
| IT | 388123-41-9P 388123-42-0P | | | | |
| | RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of tetrahydropyridine or piperidine heterocyclic derivs. and | | | | |

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their affinity for CRF receptors)

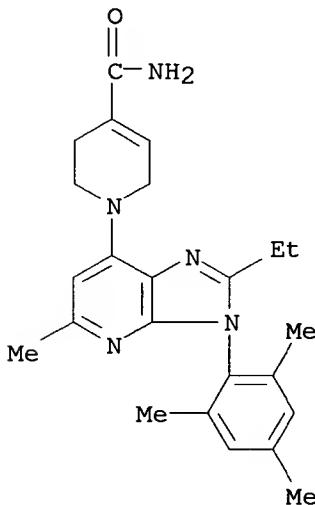
RN 388123-41-9 CAPLUS

CN 4-Pyridinecarboxamide, 1,2,3,6-tetrahydro-1-[5-methyl-3-(2,4,6-trimethylphenyl)-3H-imidazo[4,5-b]pyridin-7-yl]- (9CI) (CA INDEX NAME)



RN 388123-42-0 CAPLUS

CN 4-Pyridinecarboxamide, 1-[2-ethyl-5-methyl-3-(2,4,6-trimethylphenyl)-3H-imidazo[4,5-b]pyridin-7-yl]-1,2,3,6-tetrahydro- (9CI) (CA INDEX NAME)



RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2002 ACS

AN 1998:749412 CAPLUS

DN 130:10663

TI 2,5-Substituted arylpyrroles, compositions containing such compounds and methods of use for cytokine-mediated diseases

IN De Laszlo, Stephen E.; Liverton, Nigel J.; Ponticello, Gerald S.; Selnick, Harold G.; Mantlo, Nathan B.

PA Merck and Co., Inc., USA

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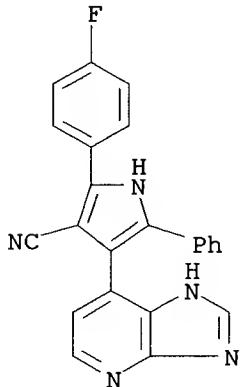
SO U.S., 48 pp.
CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|----|---|--------|----------|-----------------|----------|
| PI | US 5837719 | A | 19981117 | US 1996-694143 | 19960808 |
| OS | MARPAT 130:10663 | | | | |
| IT | 188354-63-4P | | | | |
| | RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(prepn. of 2,5-substituted arylpyrroles and treatment of cytokine-mediated diseases) | | | | |
| RN | 188354-63-4 | CAPLUS | | | |
| CN | 1H-Pyrrole-3-carbonitrile, 2-(4-fluorophenyl)-4-(1H-imidazo[4,5-b]pyridin-7-yl)-5-phenyl- (9CI) (CA INDEX NAME) | | | | |



RE.CNT 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2002 ACS

AN 1997:244347 CAPLUS

DN 126:225211

TI Preparation of triarylpyrroles as cytokine inhibitors

IN De Laszlo, Stephen E.; Mantlo, Nathan B.; Ponticello, Gerald S.; Selnick, Harold G.; Liverton, Nigel J.

PA Merck and Co., Inc., USA; De Laszlo, Stephen E.; Mantlo, Nathan B.; Ponticello, Gerald S.; Selnick, Harold G.; Liverton, Nigel J.

SO PCT Int. Appl., 126 pp.

CODEN: PIXXD2

DT Patent

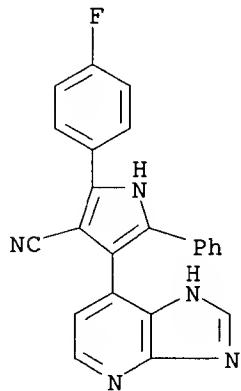
LA English

FAN.CNT 1

| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
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| PI | WO 9705878 | A1 | 19970220 | WO 1996-US12922 | 19960806 |
| | W: AL, AM, AU, AZ, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, HU, IL, IS, JP, KG, KR, KZ, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, | | | | |

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PL, RO, RU, SG, SI, SK, TJ, TM, TR, TT, UA, US, UZ, VN, AM, AZ,
 BY, KG, KZ, MD, RU, TJ, TM
 RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR,
 IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML,
 MR, NE, SN, TD, TG
 CA 2228050 AA 19970220 CA 1996-2228050 19960806
 AU 9667691 A1 19970305 AU 1996-67691 19960806
 AU 699148 B2 19981126
 EP 871444 A1 19981021 EP 1996-928099 19960806
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI
 JP 11510511 T2 19990914 JP 1996-508661 19960806
 PRAI US 1995-2094P P 19950810
 US 1996-14182P P 19960326
 GB 1996-7253 A 19960404
 GB 1996-8472 A 19960425
 WO 1996-US12922 W 19960806
 OS MARPAT 126:225211
 IT **188354-63-4P**
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. of triarylpyrroles as cytokine inhibitors)
 RN 188354-63-4 CAPLUS
 CN 1H-Pyrrole-3-carbonitrile, 2-(4-fluorophenyl)-4-(1H-imidazo[4,5-b]pyridin-7-yl)-5-phenyl- (9CI) (CA INDEX NAME)



=> log y
 COST IN U.S. DOLLARS
 FULL ESTIMATED COST

| SINCE FILE ENTRY | TOTAL SESSION |
|------------------|---------------|
| 9.56 | 150.43 |

STN INTERNATIONAL LOGOFF AT 13:28:03 ON 12 JUL 2002

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PASSWORD:

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NEWS 8 Mar 22 TRCTHERMO no longer available
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NEWS 17 Apr 22 BIOSIS Gene Names now available in TOXCENTER
NEWS 18 Apr 22 Federal Research in Progress (FEDRIP) now available
NEWS 19 Jun 03 New e-mail delivery for search results now available
NEWS 20 Jun 10 MEDLINE Reload
NEWS 21 Jun 10 PCTFULL has been reloaded
NEWS 22 Jul 02 FOREGE no longer contains STANDARDS file segment

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CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP),
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|----------------------|------------|---------|--|
| => file reg | | | |
| COST IN U.S. DOLLARS | SINCE FILE | TOTAL | |
| FULL ESTIMATED COST | ENTRY | SESSION | |
| | 0.21 | 0.21 | |

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Registry File, for complete details:
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ENTER SCREEN EXPRESSION OR (END):end

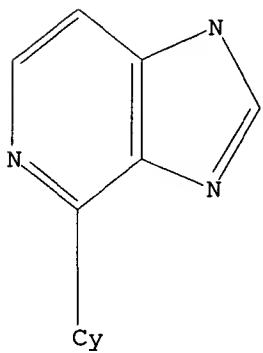
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Uploading C:\STNEXP4\QUERIES\099951592.str

L1 STRUCTURE uploaded

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L2 QUE L1

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L1 HAS NO ANSWERS
L1 STR



Structure attributes must be viewed using STN Express query preparation.

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SAMPLE SCREEN SEARCH COMPLETED - 1457 TO ITERATE

68.6% PROCESSED      1000 ITERATIONS          0 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:  ONLINE  **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:    26851 TO     31429
PROJECTED ANSWERS:        0 TO       0

L3          0 SEA SSS SAM L1

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COST IN U.S. DOLLARS           SINCE FILE          TOTAL
                                ENTRY                 SESSION
FULL ESTIMATED COST           0.38                 0.59

STN INTERNATIONAL LOGOFF AT 11:31:11 ON 12 JUL 2002
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NEWS 9 Mar 28 US Provisional Priorities searched with P in CA/CAplus and USPATFULL
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NEWS 19 Jun 03 New e-mail delivery for search results now available
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AND CURRENT DISCOVER FILE IS DATED 05 FEBRUARY 2002

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| | ENTRY | SESSION | |
| FULL ESTIMATED COST | 0.21 | 0.21 | |

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<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

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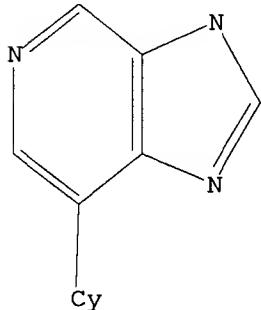
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L1 STRUCTURE uploaded

=> que L1

L2 QUE L1

=> d l1
L1 HAS NO ANSWERS
L1 STR



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